## **Delphi Consensus for Core Outcome Set for Measuring Patient Outcomes After ICU**

Instrument	Montreal Cognitive Assessment-Blind
Acronym	MoCA-Blind
Core Domain	Cognitive Sub-domain: Cognitive Screening
Area assessed (Number of questions)	Total questions: 13 Memory: 3 Attention: 4 Language: 3 Abstraction: 2 Orientation: 1
Description	A screening tool designed to detect mild cognitive dysfunction.
Versions	There are four versions of MoCA: MoCA-Full, MoCA-Blind, and MoCA-Basic. MoCA-Full is the original MoCA test. MoCA-Blind is available for use with the visually-impaired or for administration via phone. MoCA-Basic is available for use with individuals who are illiterate or with less than 5 years of formal education.
Recall Period	N/A
Scoring information	Subscores for each of the 5 sections are calculated. The total score is summed from the subscores, with a maximum score of 22. A score equal to or greater than 18 is considered normal cognition, but has not been validated thus far. Conversion from MoCA-Blind to MoCA-Full has not been validated.
Estimated time to complete	5 minutes
Administer to	Patient
Require trained administrator	Yes <sup>†</sup>
Mode of administration	In-person; via phone
Order from	http://www.mocatest.org/
Licensing Fee Fees and licensing information is effective as of September 2019, but is subject to change over time	No fee for licensing  †To access the MoCA-Blind form/manual, training certification is required from <a href="https://www.mocatest.org">www.mocatest.org</a> . Certification costs \$125 and is valid for 2 years; recertification is optional but recommended, at 50% of the initial certification cost.  Permission is required if MoCA is used for research purposes, but not for clinical or educational purposes, and can be requested via <a href="https://www.mocatest.org/permission">www.mocatest.org/permission</a> .
Equipment required	Survey form, pen, and stopwatch.
Number of published Critical Care publications using instrument (1970 – 2013) *	0
Highest COSMIN** rating (from a systematic review up to March 2015***)	No evaluation completed
Additional comments	<sup>†</sup> MoCA may be administered by anyone who understands and follows the instructions, but only a health professional with expertise in the cognitive field may interpret the results. Training in how to administer the MoCA is offered at: <a href="http://www.mocatest.org/">http://www.mocatest.org/</a>

Last updated on January 20, 2020. If you are aware of any updates required for this document, please notify us via ImproveLTO@ihmi.edu.



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The <u>full MoCA</u> has good test-retest reliability (intraclass correlation coefficient 0.87 (95% CI 0.79–0.93) and location of testing (neuro-ICU patient room vs private office) did not affect test scores. *Front Neurol*. 2019;10:734.

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<sup>\*</sup>Turnbull, A.E. et al. Outcome Measurement in ICU Survivorship Research from 1970-2013: A Scoping Review of 425 Publications. Critical Care Medicine. 2016; 44; 1267-77.

<sup>\*\*</sup> COSMIN is used to rate a study's evaluation of a survey or test's measurement properties. COSMIN does NOT rate the instrument itself, but helps readers understand if they can have confidence in the results of studies evaluating measurement properties of surveys and tests. For example, a rigorous study evaluating a test with poor measurement properties will receive a "good" COSMIN rating, while a poorly-conducted study evaluating a test with good measurement properties will receive a "poor" COSMIN rating. You must consider both the COSMIN rating and the results of studies provided when forming your opinion about that test. If more than one paper evaluated the same measurement property for a given test/survey, we present data from the paper with a better COSMIN score. COSMIN ratings were only performed for studies evaluating instruments used in ICU survivors after ICU discharge.

<sup>\*\*\*</sup>Robinson, K.A. et al. A systematic review finds limited data on measurement properties of instruments measuring outcomes in adult intensive care unit survivors. Journal of Clinical Epidemiology. 2017; 82; 37-46.